between individuals having the same or similar names.

**RECORD ACCESS PROCEDURES:**

Individuals may have access to their records by making a written request, addressed to the System Manager specified above. The envelope containing the written request must be marked ''Privacy Act Request'' or ''Freedom of Information Act Request'' or both, in the bottom left-hand corner. The letter requesting access to FCR records must state the following: (1) That the request is being made under the Privacy Act; Freedom of Information Act, or both, (2) the name, address, and signature of the requester; and (3) a detailed description of the record contents they are seeking.

**CONTESTING RECORD PROCEDURE:**

Individuals may request an amendment of a record which is not accurate, relevant, timely, or complete by writing to the System Manager at the address specified above. The envelope containing the written request must be marked ''Privacy Act Amendment Request'' or ''Freedom of Information Act Request'' or both, in the bottom left-hand corner. The letter requesting an amendment to FCR records must state the following: (1) That the request to amend the record is being made under the Privacy Act; Freedom of Information Act, or both, (2) the individual's name, address, and signature; (3) a description of the contested information; (4) the reason why the information should be amended; and (5) documentation to show that the information is inaccurate, irrelevant, untimely, or incomplete. Individuals who are contesting records must also be able to prove their identity.

**RECORD SOURCE CATEGORIES:**

Information is obtained from departments, agencies, or instrumentalities of the United States or any State.

**SYSTEM EXEMPTED FROM CERTAIN PROVISIONS OF THE ACT:**

None.

[FR Doc. 98–22581 Filed 8–21–98; 8:45 am]

**BILLING CODE 4184–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. 98C–0676]**

### Warner-Jenkinson Co., Inc.; Filing of Color Additive Petition

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that Warner-Jenkinson Co., Inc., has filed a petition proposing that the color additive regulations be amended to provide for the safe use of External D&C Violet No. 2 in coloring externally applied drug products.

**FOR FURTHER INFORMATION CONTACT:** Vivian M. Gilliam, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3167.

**SUPPLEMENTARY INFORMATION:** Under the Federal Food, Drug, and Cosmetic Act (sec. 721(d)(1) (21 U.S.C. 379e(d)(1)), notice is given that a color additive petition (CAP 8C0261) has been filed by Warner-Jenkinson Co., Inc., 107 Wade Ave., South Plainfield, NJ 07080. The petition proposes to amend the color additive regulations to provide for the safe use of External D&C Violet No. 2 in coloring externally applied drug products.

The agency has determined under 21 CFR 25.32(l) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

Dated: July 28, 1998.

**Laura M. Tarantino,**

*Acting Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.*

[FR Doc. 98–22569 Filed 8–21–98; 8:45 am]

**BILLING CODE 4160–01–F**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. 98F–0675]**

### The Dow Chemical Co.; Filing of Food Additive Petition

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that The Dow Chemical Co. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of polyethylenepolyamines as cross-linking agents for epoxy resins in coatings intended for use in contact with food.

**FOR FURTHER INFORMATION CONTACT:** Hortense S. Macon, Center for Food Safety and Applied Nutrition (HFS–205), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3086.

**SUPPLEMENTARY INFORMATION:** Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 8B4606) has been filed by The Dow Chemical Co., 2030 Dow Center, Midland, MI 48674. The petition proposes to amend the food additive regulations in § 175.300 *Resinous and polymeric coatings* (21 CFR 175.300) to provide for the safe use of polyethylenepolyamines as cross-linking agents for epoxy resins in coatings intended for use in contact with food.

The agency has determined under 21 CFR 25.32(j) that this action is of the type that does not individually or cumulatively have a significant effect on the environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

Dated: July 28, 1998.

**Laura M. Tarantino,**

*Acting Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.*

[FR Doc. 98–22570 Filed 8–21–98; 8:45 am]

**BILLING CODE 4160–01–F**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

### Arthritis Advisory Committee; Notice of Meeting

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

*Name of Committee*: Arthritis Advisory Committee.

*General Function of the Committee*: To provide advice and recommendations to the agency on FDA's regulatory issues.

*Date and Time*: The meeting will be held on September 16, 1998, 8 a.m. to 5 p.m.

*Location*: Gaithersburg Holiday Inn, Walker and Whetstone Rooms, Two Montgomery Village Ave., Gaithersburg, MD.

*Contact Person*: Gail M. Dapolito or Bill Freas, Center for Biologics Evaluation and Research (HFM–21),